Amendments to the Claims

The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (original) A method for synthesising a given peptide or its derivative which contains a proline residue or a proline derivative, at proximity to, or at, the C-terminal end of said peptide, the method comprising the steps of:
   a) synthesising on a first resin a C-terminal portion of said peptide, or its derivative, comprising at least three successive amino acid residues or their derivatives, by successive coupling of selected amino acids, small peptides or their derivatives, said first resin being suitable for the formation of peptides having a proline residue or a proline derivative positioned at, or at proximity of, the C-terminal end of said peptide;
   b) cleaving the C-terminal portion thus obtained from said first resin;
   c) reattaching said C-terminal portion to a second resin which is generally suitable for the synthesis of peptides but is unsuitable for the formation of peptides having a proline residue or a proline derivative positioned at, or at proximity of, the C-terminal end of said peptide; and
   d) coupling selected amino acids, small peptides or derivatives to the C-terminal portion to obtain said given peptide.

2. (original) The method of Claim 1 wherein said peptide is a long peptide.

3. (currently amended) The method of Claim 1 ~~or 2~~ wherein said given peptide is a chemokine having a proline residue or a proline derivative at the C-terminal or at proximity thereof.

4. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 3~~, wherein said first resin is chosen so that it does not lead to the formation of cyclic dipeptide and in particular diketopiperazine compounds.

5. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~or 4~~, wherein said step a) and/or d) is achieved by successive coupling of the predetermined amino acid residues or derivatives.

6. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 5~~, wherein said first resin for the formation of the C-terminal portion is the 2-chlorotrityl chloride resin.

7. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 6~~, wherein said second resin is a resin of the type having benzyl ester linkers.

8. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 7~~, wherein said second resin is a Wang type resin.

9. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 8~~, wherein said given peptide as up to 150 amino acid residues.

10. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 9~~, wherein the cleaving step is achieved using a mild acid treatment, for example 20% trifluoroethanol in dichloromethane.

11. (currently amended) The method of ~~any one of Claims~~ Claim 1 ~~to 10~~, wherein the C-terminal portion is fully protected so it can be attached directly onto the second resin.